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AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005

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FILE 'HOME' ENTERED AT 15:28:57 ON 30 JUN 2005

=> FIL STNGUIDE

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

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COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.06	0.27

FILE 'HOME' ENTERED AT 15:29:13 ON 30 JUN 2005

=> file agricola biosis embase caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.48

FILE 'AGRICOLA' ENTERED AT 15:29:23 ON 30 JUN 2005

FILE 'BIOSIS' ENTERED AT 15:29:23 ON 30 JUN 2005

FILE 'EMBASE' ENTERED AT 15:29:23 ON 30 JUN 2005

FILE 'CAPLUS' ENTERED AT 15:29:23 ON 30 JUN 2005

=> s phbC and phbB and phbA and transform?

=> s phbC and phbB and phbA and pseudomonas
L3 9 PHBC AND PHBB AND PHBA AND PSEUDOMONAS

=> d l3 1-9 ibib ab

L3 ANSWER 1 OF 9 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
ACCESSION NUMBER: 1999:73797 BIOSIS
DOCUMENT NUMBER: PREV199900073797
TITLE: Cloning and molecular analysis of the poly(3-hydroxybutyrate) and poly(3-hydroxybutyrate-co-3-hydroxyalkanoate) biosynthesis genes in ***Pseudomonas*** sp. strain 61-3.
AUTHOR(S): Matsusaki, Hiromi; Manji, Sumihide; Taguchi, Kazunori; Kato, Mikiya; Fukui, Toshiaki; Doi, Yoshiharu [Reprint author]
CORPORATE SOURCE: Polymer Chem. Lab., Inst. Phys. Chem. Res., 2-1 Hirosawa, Wako-shi, Saitama 351-0198, Japan
SOURCE: Journal of Bacteriology, (Dec., 1998) Vol. 180, No. 24, pp. 6459-6467. print.
CODEN: JOBAAY. ISSN: 0021-9193.
DOCUMENT TYPE: Article
LANGUAGE: English
ENTRY DATE: Entered STN: 1 Mar 1999
Last Updated on STN: 1 Mar 1999

AB Two types of polyhydroxyalkanoate (PHA) biosynthesis gene loci (phb and pha) of ***Pseudomonas*** sp. strain 61-3, which produces a blend of poly(3-hydroxybutyrate) (P(3HB)) homopolymer and a random copolymer (poly(3-hydroxybutyrate-co-3-hydroxyalkanoate) (P(3HB-co-3HA)) consisting of 3HA units of 4 to 12 carbon atoms, were cloned and analyzed at the molecular level. In the phb locus, three open reading frames encoding polyhydroxybutyrate (PHB) synthase (PhbCPs), beta-ketothiolase (PhbAPs), and NADPH-dependent acetoacetyl coenzyme A reductase (PhbBPs) were found.
The genetic organization showed a putative promoter region, followed by phbBPs-phbAPs-phbCPs. Upstream from phbBPs was found the phbRPs gene, which exhibits significant similarity to members of the AraC/XylS family of transcriptional activators. The phbRPs gene was found to be transcribed in the opposite direction from the three structural genes.
Cloning of phbRPs in a relatively high-copy vector in ***Pseudomonas*** sp. strain 61-3 elevated the levels of beta-galactosidase activity from a transcriptional phb promoter-lacZ fusion and also enhanced the 3HB fraction in the polyesters synthesized by this strain, suggesting

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=> s phbC and phbB and phbA
L4          75 PHBC AND PHBB AND PHBA

=> duplicate remove l4
DUPLICATE PREFERENCE IS 'AGRICOLA, BIOSIS, EMBASE, CAPLUS'
KEEP DUPLICATES FROM MORE THAN ONE FILE?  Y/(N):n

PROCESSING COMPLETED FOR L4
L5           51 DUPLICATE REMOVE L4 (24 DUPLICATES REMOVED)

=> d l5 1-10 ibib ab

L5   ANSWER 1 OF 51  CAPLUS  COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER:         2004:906661  CAPLUS
DOCUMENT NUMBER:          142:212974
TITLE:                    Production of polyhydroxybutyrate by
polycistronic
                          expression of bacterial genes in tobacco
plastid
AUTHOR(S):                Arai, Yuko; Shikanai, Toshiharu; Doi,
Yoshiharu;
                          Yoshida, Shigeo; Yamaguchi, Isamu; Nakashita,
Hideo
CORPORATE SOURCE:         Microbial Toxicology Laboratory, Plant
Functions
                          Laboratory, RIKEN Institute, Wako, 351-0198,
Japan
SOURCE:                   Plant and Cell Physiology (2004), 45(9), 1176-
1184
                          CODEN: PCPHA5; ISSN: 0032-0781
PUBLISHER:                Japanese Society of Plant Physiologists
DOCUMENT TYPE:            Journal
LANGUAGE:                 English
AB   Transgenic techniques are used to enhance and improve crop prodn.,
and
     their application to the prodn. of chem. resources in plants has
been
     under investigation.  To achieve this latter goal, multiple-gene
     transformation is required to improve or change plant metabolic
pathways;
     when accomplished by plant nuclear transformation, however, this
procedure
     is costly and time consuming.  Authors succeeded in the metabolic
     engineering of the tobacco plant by introducing multiple genes
within a
     bacteria-like operon into a plastid genome.  A tobacco plastid was
     transformed with a polycistron consisting of the spectinomycin
resistance
     gene and three bacterial genes for the biosynthesis of the
biodegradable
     polyester, poly[(R)-3-hydroxybutyrate] (PHB), after modification of
their
     ribosome binding sites.  DNA and RNA anal. confirmed the insertion
of the
     introduced genes into the plastid genome and their polycistronic
     expression.  As the result, the transplastomic tobacco accumulated
PHB in
     its leaves.  The introduced genes and the PHB productivity were
maternally
     inherited, avoiding genetic spread by pollen diffusion, and were
     maintained stably in the seed progeny.  Despite the low PHB
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=> s phbCAB
L6          63 PHBCAB

=> duplicate remove l6
DUPLICATE PREFERENCE IS 'AGRICOLA, BIOSIS, EMBASE, CAPLUS'
KEEP DUPLICATES FROM MORE THAN ONE FILE?  Y/(N):n

PROCESSING COMPLETED FOR L6
L7           32 DUPLICATE REMOVE L6 (31 DUPLICATES REMOVED)

=> d l7 20-32 ibib ab

L7   ANSWER 20 OF 32  BIOSIS  COPYRIGHT (c) 2005 The Thomson Corporation
on
     STN                                                    DUPLICATE 10
ACCESSION NUMBER:   2000:33419  BIOSIS
DOCUMENT NUMBER:    PREV200000033419
TITLE:              Acetate metabolism in a pta mutant of Escherichia
coli
                    W3110: Importance of maintaining acetyl coenzyme a
flux for
                    growth and survival.
AUTHOR(S):          Chang, Dong-Eun; Shin, Sooan; Rhee, Joon-Shick; Pan,
Jae-Gu
                    [Reprint author]
CORPORATE SOURCE:   Bioprocess Engineering Division, Korea Research
Institute
                    of Bioscience and Biotechnology (KRIBB), Yusong,
Taejon,
                    305-600, South Korea
SOURCE:             Journal of Bacteriology, (Nov., 1999) Vol. 181, No.
21, pp.
                    6656-6663. print.
                    CODEN: JOBAAY. ISSN: 0021-9193.
DOCUMENT TYPE:      Article
LANGUAGE:           English
ENTRY DATE:         Entered STN: 19 Jan 2000
                    Last Updated on STN: 31 Dec 2001
AB   In order to study the physiological role of acetate metabolism in
     Escherichia coli, the growth characteristics of an E. coli W3100
pta
     mutant defective in phosphotransacetylase, the first enzyme of the
acetate
     pathway, were investigated.  The pta mutant grown on glucose
minimal
     medium excreted unusual by-products such as pyruvate, D-lactate,
and
     L-glutamate instead of acetate.  In an analysis of the sequential
     consumption of amino acids by the pta mutant growing in tryptone
broth
     (TB), a brief lag between the consumption of amino acids normally
consumed
     was observed, but no such lag occurred for the wild-type strain.
The pta
     mutant was found to grow slowly on glucose, TB, or pyruvate, but it
grew
     normally on glycerol or succinate.  The defective growth and
starvation
     survival of the pta mutant were restored by the introduction of
     poly-beta-hydroxybutyrate (PHB) synthesis genes (  ***phbCAB***  )
from
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Refine Search

Search Results -

Terms	Documents
L5 and phbA and phbB and phbC	8

Database:
US Pre-Grant Publication Full-Text Database
US Patents Full-Text Database
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EPO Abstracts Database
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Derwent World Patents Index
IBM Technical Disclosure Bulletins

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Search History

DATE: Thursday, June 30, 2005 Printable Copy Create Case

Set Name side by side	Query	Hit Count	Set Name result set
DB=PGPB,USPT; PLUR=YES; OP=OR			
L6	L5 and phbA and phbB and phbC	8	L6
L5	huisman-Gjalt-$.in.	15	L5
DB=USPT; PLUR=YES; OP=OR			
L4	L1 and PHA and bacteria.clm.	12	L4
L3	L1 and PHA and bacteria	39	L3
DB=PGPB,USPT,USOC,EPAB,JPAB,DWPI; PLUR=YES; OP=OR			
L2	L1 and PHA	57	L2
L1	phbC and phbB and phbA and bacteria	61	L1

END OF SEARCH HISTORY